Amendments to the Claims:

The following listing of claims replaces all prior versions and listings of the claims in this application.

Listing of the Claims

1. (Currently amended) A method for proliferating cardiomyocytes comprising: introducing nucleotide sequences coding for a D-type cyclin gene and a cyclin dependent kinase gene directly into the cardiomyocytes using a <u>viral</u> vector <u>and expressing said nucleotide sequences in said cardiomyocytes</u>, and cultivating or holding said cardiomyocytes,

wherein said cyclin gene is a gene coding for cyclin D1, D2 or D3, wherein said cyclin dependent kinase gene is a gene coding for CDK4 or CDK6, and wherein a nucleotide sequence coding for a nuclear localization signal is attached to at least one of said cyclin gene or said cyclin dependent kinase gene.

2. (Currently amended) A method for proliferating cardiomyocytes comprising: introducing nucleotide sequences coding for a D-type cyclin gene and a cyclin dependent kinase gene into cardiomyocytes in vitro using a viral vector and expressing said nucleotide sequences in said cardiomyocytes, and then cultivating said cardiomyocytes, or introducing each of said genes directly to cardiomyocytes in vivo using a viral vector and expressing said genes in said cardiomyocytes,

wherein said cyclin gene is a gene coding for cyclin D1, D2 or D3, wherein said cyclin dependent kinase gene a gene coding for is CDK4 or CDK6, and wherein a nucleotide sequence coding for a nuclear localization signal is attached to at least one of said cyclin gene or said cyclin dependent kinase gene.

- 3. (Canceled)
- 4. (Canceled)
- 5. (Canceled)
- 6. (Currently amended) The method of claim 2, wherein said eyelin-gene and said eyelin-dependent kinase gene are transferred to the cardiomyoeytes using viral vector is an adenovirus vector.

- 7. (Withdrawn) A recombinant vector comprising a cyclin gene comprising a nucleotide sequence coding for a nuclear localization signal.
- 8. (Withdrawn) A recombinant vector comprising a cyclin gene and a cyclin dependent kinase gene, wherein at least one of said genes is attached with a nucleotide sequence coding for a nuclear localization signal.
- 9. (Canceled)
- 10. (Canceled)
- 11. (Canceled)
- 12. (Canceled)
- 13. (Canceled)
- 14. (Canceled)
- 15. (Canceled)
- 16. (Previously presented) The method of claim 2, wherein said genes comprising said nucleotide sequences are introduced to the cardiomyocytes *in vitro*, and cultivating said cardiomyocytes.
- 17. (Previously presented) The method of claim 2, wherein said genes comprising said nucleotide sequences are introduced to the cardiomyocytes *in vivo*.
- 18. (Previously presented) The method of claim 1 or 2, wherein said cyclin activates CDK4.
- 19. (Previously presented) The method of claim 1 or 2, wherein said cyclin activates CDK6

- 20. (Previously presented) The method of claim 2, wherein said cyclin is D1.
- 21. (Previously presented) The method of claim 1, wherein the cyclin is D2 or D3.
- 22. (Previously presented) The method of claim 2, wherein the cyclin is D2 or D3.
- 23. (Previously presented) The method of claim 1, wherein the cyclin dependent kinase is CDK4.
- 24. (Previously presented) The method of claim 1, wherein the D-type cyclin is D1.
- 25. (Previously presented) The method of claim 16, wherein the cyclin dependent kinase is CDK4.
- 26. (Previously presented) The method of claim 16, wherein the D-type cyclin is D1.
- 27. (Previously presented) The method of claim 16, wherein the cyclin dependent kinase is CDK4 and the D-type cyclin is D1.
- 28. (Previously presented) The method of claim 17, wherein the cyclin dependent kinase is CDK4.
- 29. (Previously presented) The method of claim 17, wherein the D-type cyclin is D1.
- 30. (Previously presented) The method of claim 17, wherein the cyclin dependent kinase is CDK4 and the D-type cyclin is D1.
- 31. (Canceled)
- 32. (Canceled)

- 33. (Canceled)
- 34. (Canceled)
- 35. (Canceled)
- 36. (Canceled)
- 37. (Currently amended) A method for proliferating cardiomyocytes *in vitro* comprising: introducing nucleotide sequences coding for a D-type cyclin and a recombinant cyclin dependent kinase gene directly into the cardiomyocytes using a <u>viral</u> vector <u>and expressing said nucleotide</u> sequences in said cardiomyocytes, and cultivating or holding said cardiomyocytes,

wherein said cyclin gene is a gene coding for cyclin D1, D2 or D3,
wherein said cyclin dependent kinase gene is a gene coding for CDK4 or CDK6, and
wherein a nucleotide sequence coding for a nuclear localization signal is attached to at least
one of said cyclin gene or said cyclin dependent kinase gene.

38. (Currently amended) A method for proliferating cardiomyocytes *in vivo* comprising: introducing nucleotide sequences coding for a D-type cyclin gene and a cyclin dependent kinase gene directly to cardiomyocytes *in vivo* using a viral vector and expressing said nucleotide sequences in said cardiomyocytes,

wherein said cyclin is cyclin D1, D2 or D3,

wherein said cyclin dependent kinase is CDK4 or CDK6, and

wherein a nucleotide sequence coding for a nuclear localization signal is attached to at least one of said cyclin gene or said cyclin dependent kinase gene.

39. (Currently amended) The method of claim 1, wherein said eyelin gene and said eyelin dependent kinase gene are transferred to the cardiomyocytes using viral vector is an adenovirus vector.